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                 Web Page for STN Seminar Schedule - N. America
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        JUL 02
                LMEDLINE coverage updated
        JUL 02
NEWS
                SCISEARCH enhanced with complete author names
NEWS
        JUL 02
                CHEMCATS accession numbers revised
        JUL 02 CA/CAplus enhanced with utility model patents from China
NEWS
NEWS 6 JUL 16 CAplus enhanced with French and German abstracts
        JUL 18 CA/CAplus patent coverage enhanced
NEWS 7
NEWS 8 JUL 26 USPATFULL/USPAT2 enhanced with IPC reclassification
NEWS 9 JUL 30 USGENE now available on STN
NEWS 10 AUG 06 CAS REGISTRY enhanced with new experimental property tags
NEWS 11 AUG 06 FSTA enhanced with new thesaurus edition
NEWS 12
        AUG 13 CA/Caplus enhanced with additional kind codes for granted
                 patents
NEWS 13
        AUG 20
                 CA/CAplus enhanced with CAS indexing in pre-1907 records
NEWS 14
       AUG 27
                 Full-text patent databases enhanced with predefined
                 patent family display formats from INPADOCDB
NEWS 15
        AUG 27
                USPATOLD now available on STN
                CAS REGISTRY enhanced with additional experimental
NEWS 16 AUG 28
                 spectral property data
NEWS 17
        SEP 07
                 STN AnaVist, Version 2.0, now available with Derwent
                 World Patents Index
         SEP 13
                 FORIS renamed to SOFIS
NEWS 18
NEWS 19
         SEP 13
                 INPADOCDB enhanced with monthly SDI frequency
NEWS 20
        SEP 17
                 CA/CAplus enhanced with printed CA page images from
                 1967-1998
NEWS 21
        SEP 17
                 CAplus coverage extended to include traditional medicine
                 patents
NEWS 22
         SEP 24
                 EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS 23
         OCT 02
                 CA/CAplus enhanced with pre-1907 records from Chemisches
                 Zentralblatt
NEWS 24
        OCT 19
                BEILSTEIN updated with new compounds
NEWS EXPRESS
             19 SEPTEMBER 2007: CURRENT WINDOWS VERSION IS V8.2,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 19 SEPTEMBER 2007.
NEWS HOURS
              STN Operating Hours Plus Help Desk Availability
NEWS LOGIN
              Welcome Banner and News Items
NEWS IPC8
              For general information regarding STN implementation of IPC 8
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=> file registry
COST IN U.S. DOLLARS

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FULL ESTIMATED COST

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STRUCTURE FILE UPDATES: 19 OCT 2007 HIGHEST RN 951118-42-6 DICTIONARY FILE UPDATES: 19 OCT 2007 HIGHEST RN 951118-42-6

New CAS Information Use Policies, enter HELP USAGETERMS for details.

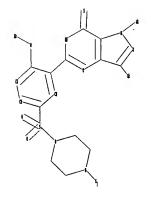
TSCA INFORMATION NOW CURRENT THROUGH June 29, 2007

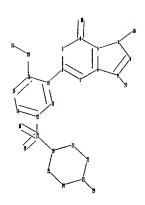
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http://www.cas.org/support/stngen/stndoc/properties.html

Uploading C:\Program Files\Stnexp\Queries\10 series\10583335\10583335a.str





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chain nodes :
10 17 18 19 26 27 28 29 31
ring nodes :
1 2 3 4 5 6 7 8 9 11 12 13 14 15 16 20 21 22 23 24 25
chain bonds :
2-11 4-10 7-28 9-27 13-17 16-26 17-18 17-19 17-20 23-29 26-31
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9 11-12 11-16 12-13 13-14 14-15
15-16 20-21 20-25 21-22 22-23 23-24 24-25
exact/norm bonds :
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exact bonds :
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normalized bonds :
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G1:H,Ak,Cb

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS 20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom 26:CLASS 27:CLASS 28:CLASS 29:CLASS 31:CLASS

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 13:43:09 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 89 TO ITERATE

100.0% PROCESSED 89 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 1214 TO 2346

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 13:43:13 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 1639 TO ITERATE

100.0% PROCESSED 1639 ITERATIONS 25 ANSWERS

SEARCH TIME: 00.00.01

L3 25 SEA SSS FUL L1

=> d scan

25 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN Piperazine, 1-[[3-(1-ethyl-4,7-dihydro-3-propyl-7-thioxo-1H-pyrazolo[4,3-dlpyrimidin-5-yl)-4-propoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,5S)-rel-(9CI)
C25 H36 N6 O3 S2

Relative stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):3

L3 25 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN

Piperazine,

1-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-dipyrimidin-5-yl)-4-ethoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,5S)-rel-(9CI)

MF C23 H32 N6 O3 S2

CI COM

Relative stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 25 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Piperazine,
1-[{3-{47-7d.hydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo{4,3-d|pyrimidin-5-yl)-4-propoxyphenyl}sulfonyl}-3,5-dimethyl-, (3R,5S)-rel-(9CI)
MF C24 H34 N6 03 S2

Relative stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 25 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN
IN Piperazine,
1-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-(3-methylbutoxy)phenyl]sulfonyl]-4-ethylMF C26 H38 N6 O3 S2

**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT **

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> file caplus
COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 173.00 173.21

FILE 'CAPLUS' ENTERED AT 13:43:52 ON 22 OCT 2007
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=> s 13

L4 4 L3

=> d l4 1-4 ibib abs hitstr

L4 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS ON STN ACCESSION NUMBER: 2005:570896 CAPLUS DOCUMENT NUMBER: 143:97390

NATIONAL ACTIONS

143:97390

Preparation of pyrazolopyrimidinethione derivatives for treatment of impotence
Li, Shuxin: Ren, Jianping: Zhao, Yanjin; Lv, Qiujun; Quo, Jinhua
The Institute of Radiation Medicine, Academy of Millitary Medical Sciences Pla, Peop. Rep. China PCT Int. Appl., 34 pp.
CODEN: PIXXD2
Patent
Chinese
1 TITLE: INVENTOR (S):

PATENT ASSIGNEE(S):

DOCUMENT TYPE: LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

| | PATENT NO. | | | | | | | | | | | | | | | | | | |
|----------------|--------------------------|------|-------|-------|-----|-------------|-----|-----|-----|------------------|-------|-------|----------|------|------|-----|------|-----|--|
| | WO 2005058899 | | | | | | | | | | | | | | | | | | |
| | | | | | | A1 20050630 | | | | WO 2 | 004- | | 20041118 | | | | | | |
| | | W: | ΑE, | AG, | AL, | AM, | AT, | ΑU, | AZ, | BA, | BB, | BG, | BR, | BW, | BY, | ΒZ, | CA, | CH, | |
| | | | CN, | co. | CR, | CU, | CZ. | DE. | DK, | DM, | DZ, | EC, | EE, | EG, | ES, | FI, | GB, | GD, | |
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| | CN 1629163 EP 1695976 | | | | | | | | | CN 2003-10118481 | | | | | | | | | |
| | | | | | | A1 20060830 | | | | EP 2 | 004- | | 20041118 | | | | | | |
| | | R: | AT, | BE, | CH, | DE, | DK, | ES, | FR, | GB, | GR, | IT, | LI, | LU, | NL, | SE, | MC, | PT, | |
| | | | IE. | SI. | FI. | RO. | CY. | TR. | BG. | CZ. | EE. | HU, | PL. | SK. | IS | | | | |
| IN 2006MN00737 | | | | | | | | | | | | | | 2 | 0060 | 623 | | | |
| US 2007219220 | | | | | | | | | | | | | | | | | | | |
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OTHER SOURCE(S):

CASREACT 143:97390; MARPAT 143:97390

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

IT 856190-48-2P 856190-49-3P 856190-50-6P 856190-51-7P 856190-56-2P RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of pyrazolopyrimidinethione deriva. for treatment of impotence)
RN 856190-48-2 CAPLUS

impotence)
impotence;
RN 856190-48-2 CAPLUS
CN Piperazine,
1-{(3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo(4,3-dipyrimidin-5-yl)-4-methoxyphenyl]sulfonyl}-3,5-dimethyl-, (3R,55)-rel-(9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 856190-49-3 CAPLUS
CN Piperazine,
1-[(3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo(4,3-dlpyrimidin-5-yl)-4-propoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,5s)-rel-(9CI) (CA INDEX NAME)

Relative stereochemistry.

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS OR STN (Continued)

AB Title compds. represented by the formula I [wherein R1-R3 = independently ([cyclo]alkoxy]alkyl, alkenyl or aryl; R4 = alkyl, alkenyl, (cyclo]alkoxy, aryl; R6 = H, (cyclo]alkox, aryl; R6 = H, (cyclo]alkyl, alkenyl, alkylcarbonyl; and pharmaceutically acceptable salts or solvates thereof) were prepared for treatment of impotence. For example, II was given in a multi-step synthesis starting from 4-amino-1-ethyl-3- propylpyrazole-5-carboxamide. I showed enhanced erectile response in rats

rats similar to that of Sildenafil. Thus, I and their pharmaceutical compnsare useful for the treatment of impotence and sexlessness, having high selectivity over PDE V, long action time, less side reactions, and no

side effects of blood pressure decreasing and heart rate increasing. $856190\hbox{--}47\hbox{--}1P$

IT 856190-47-1P
RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); SPN (Synthetic preparation); TRU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of pyrazolopyrimidinethione derivs. for treatment of impotence)

856190-47-1 CAPLUS

RN 83613U-41-1 CARDON
Piperazine,
1-[(3-(4,7-dih)dro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-dipyelmidin-5-yl)-4-ethoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,5s)-rel(9CI) (CA INDEX NAME)

Relative stereochemistry.

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

856190-50-6 CAPLUS
Piperazine, 1-[{3-(1-ethyl-4,7-dihydro-3-propyl-7-thioxo-1H-pyrazolo[4,3-d|pyrimidin-5-yl)-4-methoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,5S)-rel-(9CI) (CA INDEX NAME)

Relative stereochemistry.

856190-51-7 CAPLUS
Piperazine, 1-[[3-(1-ethyl-4,7-dihydro-3-propyl-7-thioxo-1H-pyrazolo[4,3-d)pyrimidin-5-yl)-4-propoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,58)-rel-(9CI) (CA INDEX NAME)

Relative stereochemistry.

856190-56-2 CAPLUS

RN 856190-56-2 CAPLUS
CN Piperazine,
1-{[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued) djpyrimidin-5-yl)-4-ethoxyphenyl]sulfonyl]-3,5-dimethyl-, (3R,5S)-rel-,2-hydroxy-1,2,3-propanetricarboxylate (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 856190-47-1 CMF C23 H32 N6 O3 S2

Relative stereochemistry.

CM 2

CRN 77-92-9 C6 H8 O7

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

RN 479073-79-5 CAPLUS
CN Piperazine,
[-[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl]-4-ethoxyphenyl]sulfonyl]-4-methyl- (9CI)
NAME)

RN 479073-80-8 CAPLUS
CN Piperazine,
1-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-ethoxyphenyl]sulfonyl]-4-ethyl- (9CI) (CA INDEX

RN 479073-86-4 CAPLUS
CN Piperazine,
1-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl]-4-ethoxyphenyl]aulfonyl]- (9CI) (CA INDEX NAME)

L4 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS ON STN ACCESSION NUMBER: 2005:565628 CAPLUS DOCUMENT NUMBER: 143:211579
TITLE: Low-energy 2011-1

143:211579
Low-energy collision-induced dissociation of sildenafil thiono analogues: Gas-phase intramolecular nucleophilic substitution through ion-neutral complexes between a cationic substrate and a thione-containing neutral nucleophile Lee, Jaeick; Yoo, Hye Hyan; Kang, Min-Yung; Kim, Dong-Hyun Bioanalysis and Biotransformation Research Center, Korea Institute of Science and Technology, Seoul, S. Korea

AUTHOR (S):

CORPORATE SOURCE:

Korea Rapid Communications in Mass Spectrometry (2005), 19(12), 1767-1770
CODEN: RCMSEF; ISSN: 0951-4198
John Wiley & Sons Ltd.

PUBLISHER: DOCUMENT TYPE:

Journal LANGUAGE:

WENT TYPE: JOURNAL JOU

IT

dipyrim.din-3-11-4-e-insyphenylistitonyli-4-(methylipiperazine) were reported.
479073-72-8 479073-74-0 479073-79-5
479073-80-8 479073-86-4
RL: PRP (Properties)
(study of low energy collision-induced dissociation of sildenafil and

thioxo analogs and study of gas-phase intramol. nucleophilic substitution through ion-neutral complexes between cationic substrate and thione-containing neutral nucleophile)

RN 479073-72-8 CAPLUS

CN Piperazine,
1-[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-methoxyphenyl]sulfonyl]-4-methyl- (9CI) (CA INDEX NAME) its

RN 479073-74-0 CAPLUS
CN Piperazine,
[-[3-4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-methoxyphenyl]sulfonyl)-4-ethyl- (9CI) (CA INDEX NAME)

ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

REFERENCE COUNT: THIS

11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR

FORMAT

RECORD. ALL CITATIONS AVAILABLE IN THE RE

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ACCESSION NUMBER: 2004:524375 CAPLUS
DOCUMENT NUMBER: 141:420374

TITLE: Effects of a new selective phosphodiesterase type 5 inhibitor, KJH-1002, on the relexation of rabbit corpus cavernosum tissue

AUTHOR(S): Cho, Eun Young: Chung, Sung-Hyun; Kim, Joong Hyup; Kim, Dong-Kyun; Jin, Changbase

CORPORATE SOURCE: Bloanlysis (# Biotransformation Research Center, Korea

Institute of Science and Technology, Seoul, 130-650, S. Korea

SOURCE: Journal of Applied Pharmacology (2003), 11(4),

232-237

CODEN: JOAPA6; ISSN: 1225-6110

PUBLISHER: Korean Society of Applied Pharmacology

DOCUMENT TYPE: Journal

LANGGUAGE: English

AB The present study examined functional effects of a new selective phosphodiesterase type 5 inhibitor, 1-(4-ethoxy-3-(6,7-dihydro-1-methyl-7-thioxo-3-propyl-1H-pyrazolo[4,3]pyrimidin-5-yl)phenylsulfonyl]-4-Me piperazine (KJH-1002), in the isolated rabbit corpus cavernosum (RCC). Relaxing effects of KJH-1002 were also compared with those of sildensfil, which is currently used as an oral therapy for penile erectile dysfunction. In the isolated RCC precentracted with phenylephrine, both KJH-1002 and sildensfil in the concentration-dependent manner. In the sodium nitroprusside relaxation, the Ic50 values, concns. of SNP required to produce a 50% relaxation, the Ic50 values, concns. of SNP required to produce a 50% relaxation, the Ic50 values, concns. of SNP required to produce a 50% relaxation of the phenylephrine-induced contraction, were significantly decreased to the similar extent by treatments with KJH-1002 and sildensfil. The results suggest that a new selective phosphodiesterase type 5 inhibitor, KJH-1002, has an augmentative effect on penile erection comparable to that of sildensfil and can be useful for the treatment of erectile dysfunction.

IT 479073-79-5

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(KJH-1002) phosphodiesterase type-5 inhibitor produced relaxing effect on rabbit corpus cavernosum and showed augmentative effect on penile ere
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L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS ON STN
ACCESSION NUMBER:
DOCUMENT NUMBER:
138:55978
128:55978
171LE:
Preparation of novel pyrazolopyrimidinethiones as phosphodiesterase V inhibitors for treating erectile dysfunction
NVENTOR(S):
Kim, Joong-Hyup: Kim, Youseung: Choi, Kyung II; Kim, Dong Hyun: Nam, Ghilsoo; Seo, Jae Hong
Korea Institute of Science and Technology, S. Korea Pet Tint. Appl., 34 pp.
CODEN: PIXXD2
PATENT INFORMATION:
PATENT INFORMATION:

| | PAT | TENT | NO. | | KIN | D | DATE | | | APP | LICAT | DATE | | | | | | | | |
|-----------------|-----|---------------|------|-----|-----|-----|-------------|------|---------------|-----|-------|-------|------|----------|-----|-----|------|-----|--|--|
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| | WO | WO 2002102802 | | | | | A1 20021227 | | | , | WO | 2002- | | 20020614 | | | | | | |
| | | W: | ΑE, | AG, | AL, | AM, | AT, | ΑU, | ΑZ, | BA, | BB | , BG, | BR, | BY, | BZ, | CA, | CH, | CN, | | |
| | | | co, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ, | EC | , EE, | ES, | FI, | GB, | GD, | GE, | GH, | | |
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| | | | BF. | BJ. | CF. | CG. | CI. | CM. | GA. | GN. | GO | , GW, | ML. | MR. | NE. | SN. | TD. | TG | | |
| | | | | | | | | | KR 2001-33382 | | | | | | | | | | | |
| AU 2002315822 . | | | | | | | | | | | | | | | | | | | | |
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| | | | | | | | | | | | | TR | | | | | | | | |
| | JР | 2005 | 5055 | 09 | | T | | 2005 | 0224 | | JP | 2003- | 5062 | 75 | | 2 | 0020 | 614 | | |
| | | | | | | | | | | | | 2003- | | | | | | | | |
| PRIOR | | | | | | | | | | | | 2001- | | | | | | | | |
| | | | | | | | | | | | | | | - | | | | | | |
| | | | | | | | | | | , | wo | 2002- | KR11 | 26 | | w 2 | 0020 | 614 | | |

OTHER SOURCE(S): CASREACT 138:55978; MARPAT 138:55978

L4 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

REFERENCE COUNT: THIS 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)
AB The title compds. [I: R1, R2 = H, alkyl, cycloalkyl; R3 = alkyl, cycloalkyl or alkenyl which is unsubstituted or substituted; X = 0, NR4; R4 = H, alkyl, cycloalkyl or alkenyl which is unsubstituted or substituted with OH or alkoxyl which exhibit higher inhibitory activities against phosphodiesterase V as well as lower inhibitory activities against phosphodiesterase isoenzymes I, III and VI (biol. data given) and therefore are useful for the treatment of erectile dysfunction, were prepared E.G., a 3-step synthesis of I [R1 = Me; R2 = Pr; R3 = Et; X = NMe], starting from 5-(2-ethoxyphenyl)-1-methyl-3-propyl-1,6-dihydropyrazolo[4,3-d]pyrimidin-7-one, which showed IC50 of 0.59 nM against PDE V, was given.

IT 479073-72-8P 479073-80-8P 479073-86-6P 479073-79-8P 479073-89-8P 479073-89-8P 479073-89-8P 479073-89-8P 479073-89-8P 479073-97-P 479073-98-8P 479073-98-8P 479073-97-P 479073-97-P 479073-98-8P 479074-02-P RI: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of novel pyrazolopyrimidinethiones as PDE5 inhibitors for treating erectile dysfunction)

RN 479073-72-8 CAPLUS

Piperazine,
1-[(3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-dipyrimidin-5-yl)-4-methoxyphenyl]sulfonyl]-4-methyl- (9CI) (CA INDEX NAME)

479073-74-0 CAPLUS
Piperazine,
3-(4,7-dihydro-1-methy1-3-propy1-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-y1)-4-methoxyphenyl]sulfonyl]-4-ethyl- (9CI) (CA INDEX NAMP)

L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)
RN 479073-76-2 CAPLUS
CN 1-Piperazineethanol, 4-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-methoxyphenyl]sulfonyl]- (9CI) (CA INDEX NAME)

RN 479073-79-5 CAPLUS
CN Piperazine,
[-[3-(4,7-dihydro-1-methy1-3-propy1-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-y1)-4-ethoxyphenyl]sulfonyl]-4-methyl- (9CI) (CA INDEX NAME)

RN 479073-80-8 CAPLUS
CN Piperazine,
1-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-ethoxyphenyl]sulfonyl]-4-ethyl- (9CI) (CA INDEX NAME)

479073-82-0 CAPLUS
1-Piperazineethanol, 4-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo(4,3-d]pyrimidin-5-yl)-4-ethoxyphenyl]sulfonyl]- (9CI) (CA INDEX NAME)

ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

RN 479073-90-0 CAPLUS
CN 1-Piperazineethanol, 4-[[3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo(4,3-d]pyrimidin-5-yl)-4-propoxyphenyl]sulfonyl]- (9CI) (CA INDEX

NAME)

479073-92-2 CAPLUS
Piperazine, 1-[(4-butoxy-3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)phenyl)aulfonyl]-4-methyl- (9CI) (CA INDEX NAME)

479073-93-3 CAPLUS
Piperazine, 1-[{4-butoxy-3-{4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)phenyl]sulfonyl]-4-ethyl- (9CI) (CA INDEX NAME)

ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

RN 479073-86-4 CAPLUS CN Piperazine, 1-[{3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-dipyrimidin-5-yl)-4-ethoxyphenyl]sulfonyl]- (9CI) (CA INDEX NAME)

RN 479073-87-5 CAPLUS

Piperazine,
[[3-4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-propoxyphenyl]sulfonyl]-4-methyl- (9CI) (CA INDEX NAME)

RN 479073-88-6 CAPLUS
CN Piperazine,
1-[[3-(4,7-dih)dro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl]-4-propoxyphenyl]sulfonyl]-4-ethyl- {9Cl} (CA INDEX NAME)

ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

479073-94-4 CAPLUS
1-Piperazineethanol, 4-[{4-butoxy-3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)phenyl}sulfonyl]- (9CI) (CA

RN 479073-96-6 CAPLUS
CN Piperazine,
-[[3-[4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl]-4-(2-methylpropoxy)phenyl]sulfonyl]-4-methyl- (9CI)

RN 479073-97-7 CAPLUS ,
CN Piperazine,
-[(3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-(2-methylpropoxy)phenyl]sulfonyl]-4-ethyl- (9CI) (CA INDEX NAME)

L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

479073-98-8 CAPLUS
1-Piperazineethanol, 4-[{3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo(4,3-d)pyrimidin-5-yl)-4-(2-methylpropoxy)phenyl)sulfonyl)- (9CI) (CA INDEX NAME)

RN 479074-00-5 CAPLUS
CN Piperazine,
1-[[3-(4,7-d.hydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimldin-5-yl)-4-(3-methylbutoxy)phenyl]sulfonyl]-4-methyl- (9CI) (CA INDEX NAME)

RN 479074-01-6 CAPLUS

Piperazine,

1-([3-(4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo{4,3-dipyrimidin-5-yl)-4-(3-methylbutoxy)phenyl]aulfonyl]-4-ethyl- (9CI) (CA INDEX NAME)

L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

479074-02-7 CAPLUS 1-Piperazineethanol, 4-[[3-[4,7-dihydro-1-methyl-3-propyl-7-thioxo-1H-pyrazolo[4,3-d]pyrimidin-5-yl)-4-[3-methylbutoxy)phenyl]sulfonyl]- (9CI) (CA INDEX NAME)

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